PATENT NO. : 7,214,690 B2

APPLICATION NO.: 10/080503

Page 1 of 11

DATED INVENTOR(S)

: May 8, 2007 : Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

IN THE TITLE PAGES:

In Item [56] References Cited, in OTHER PUBLICATIONS: in Venturoli et al., please replace "Prospectiove" with --Prospective-

IN THE SPECIFICATION:

At column 5, line 20, please replace structure

with the following structure: --

At column 9, line 34, please replace "A R⁹" with --R⁹--At column 31, line 56-67, please replace structure

with the following structure: --

PATENT NO.

: 7,214,690 B2

Page 2 of 11

APPLICATION NO.: 10/080503

DATED

: May 8, 2007

INVENTOR(S)

: Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

IN THE CLAIMS:

Please replace Claims 1, 10, 24, 40, 57, and 58 with the following Claims:

1. A compound having the formula:

Column 79, line 40 thry Column 81, line 37 Shovid read

(I)

wherein:

R¹ is selected from the group consisting of hydrogen, F, Cl, Br, I, NO₂, OR⁹, NR¹⁰R¹¹, S(O)_nR⁹, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C1-C8 heteroalkyl, optionally substituted C₃-C₈ cycloalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted C2-C8 alkynyl and optionally substituted C₂-C₈ alkenyl;

R² is selected from the group consisting of hydrogen, F, Cl, Br, I, CF₃, CF₂Cl, CF₂H, CFH₂, CF₂OR⁹, CH₂OR⁹, OR⁹, S(O)_nR⁹, NR¹⁰R¹¹, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted C3-C8 cycloalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted C_2 – C_8 alkynyl and optionally substituted C_2 – C_8 alkenyl;

R³ and R⁴ each independently is selected from the group consisting of hydrogen, OR⁹, S(O)_nR⁹, NR¹⁰R¹¹, C(Y)OR¹¹, CNR¹⁰R¹¹, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted C3-C8 cycloalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted C₂-C₈ alkynyl and optionally substituted C₂-C₈ alkenyl;

PATENT NO.

: 7,214,690 B2

Page 3 of 11

APPLICATION NO.: 10/080503

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: May 8, 2007

INVENTOR(S)

: Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

R⁵ and R⁶ each independently is selected from the group consisting of hydrogen, CF₃, CF₂Cl, CF₂H, CFH₂, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted C3-C8 cycloalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted C2-C8 alkynyl and optionally substituted C2-C8 alkenyl;

R⁷ is selected from the group consisting of hydrogen, F, Cl, Br, I, optionally substituted C1-C8 alkyl, optionally substituted C1-C8 haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl, OR⁹,

S(O)_nR⁹, NR¹⁰R¹¹, C(Y)OR¹¹ and C(Y)NR¹⁰R¹¹;

R⁸ is selected from the group consisting of hydrogen, F, Cl, Br, I, optionally substituted C_1 – C_8 alkyl, optionally substituted C_1 – C_8 haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl, OR⁹, $S(O)_n R^9$, $NR^{10}R^{11}$, $C(Y)OR^{11}$ and $C(Y)NR^{10}R^{11}$;

R9 is selected from the group consisting of hydrogen, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl, and optionally

substituted arylalkyl;

R¹⁰ is selected from the group consisting of hydrogen, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted arylalkyl, CO_2R^{12} , $C(O)R^{12}$, SO_2R^{12} and $S(O)R^{12}$; R^{11} and R^{12} each independently is selected from the group consisting of

hydrogen, optionally substituted C_1 – C_8 alkyl, optionally substituted C_1 – C_8 haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally

substituted heteroaryl and optionally substituted arvlalkyl:

R¹³ is selected from the group consisting of optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted C2-C8 alkenyl, optionally substituted C2-C8 alkynyl, optionally substituted C3-C8 cycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted arylalkyl and optionally substituted heteroarylalkyl;

m is selected from the group consisting of 0, 1 and 2; n is selected from the group consisting of 0, 1 and 2;

W is selected from the group consisting of NH, N{R¹³}, N{C(Y)R¹¹} and $N{SO_2R^{11}};$

X is O:

Z is selected from the group consisting of NH, $N\{R^{11}\}$, $N\{C(Y)R^{11}\}$, $N{SO_2R^{12}}$ and $N{S(O)R^{12}}$; and

Y is O:

and pharmaceutically acceptable salts thereof; wherein:

PATENT NO.

: 7,214,690 B2

Page 4 of 11

APPLICATION NO.: 10/080503

DATED

: May 8, 2007

INVENTOR(S)

: Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

the substituents of an optionally substituted group comprise one or more substituents independently selected from among alkyl, alkenyl, alkynyl, heteroalkyl, haloalkyl, haloalkenyl, haloalkynyl, cycloalkyl, aryl, heteroaryl, arylalkyl, heteroarylalkyl, alkoxy, aryloxy, haloalkoxy, amino, alkylamino, dialkylamino, alkylthio, arylthio, heteroarylthio, oxo, carboxyester, carboxamido, acyloxy, hydrogen, F, Cl, Br, I, CN, NO₂, NH₂, N₃, NHCH₃, N(CH₃)₂, SH, SCH₃, OH, OCH₃, OCF₃, CH₃, CF₃, C(O)CH₃, CO₂CH₃, CO₂H, C(O)NH₂, OR⁹, SR⁹, NR¹⁰R¹¹, CF₂CF₃, CH₂CH₂F and CH2CF3. Column 82, lines 6-54, Claim 10 should read

10. The compound of claim 1, wherein:

R¹ is selected from the group consisting of hydrogen, F, Cl, Br, I, NO₂, OR⁹, $NR^{10}R^{11},\,S(O)_nR^9,\,C_1-C_8\,\,alkyl,\,C_1-C_8\,\,haloalkyl,\,C_1-C_8\,\,heteroalkyl,\,C_3-C_8\,\,cycloalkyl,$

aryl, arylalkyl, heteroaryl, C2-C8 alkynyl and C2-C8 alkenyl;

R² is selected from the group consisting of hydrogen, F, Cl, Br, I, CF₃, CF₂Cl, CF_2H , CFH_2 , CF_2OR^9 , CH_2OR^9 , OR^9 , $S(O)_nR^9$, $NR^{10}R^{11}$, C_1 – C_8 alkyl, C_1 – C_8 haloalkyl, C₁-C₈ heteroalkyl, C₃-C₈ cycloalkyl, aryl, arylalkyl, heteroaryl, C₂-C₈ alkynyl and C₂-C₈ alkenyl;

R³ and R⁴ each independently is selected from the group consisting of hydrogen, OR⁹, S(O)_nR⁹, NR¹⁰R¹¹, C(Y)OR¹¹, C(Y)NR¹⁰R¹¹, C₁-C₈ alkyl, C₁-C₈ haloalkyl, C₁-C₈ heteroalkyl, C₃-C₈ cycloalkyl, aryl, arylalkyl, heteroaryl, C₂-C₈ alkynyl and C_2 – C_8 alkenyl;

R⁵ and R⁶ each independently is selected from the group consisting of hydrogen, CF₃, CF₂Cl, CF₂H, CFH₂, C₁-C₈ alkyl, C₁-C₈ haloalkyl, C₁-C₈ heteroalkyl, C₃-C₈ cycloalkyl, aryl, arylalkyl, heteroaryl, C₂-C₈ alkynyl and C₂-C₈ alkenyl;

R⁷ is selected from the group consisting of hydrogen, F, Cl, Br, I, C₁-C₈ alkyl, C₁-C₈ haloalkyl, C₁-C₈ heteroalkyl, aryl, heteroaryl, OR⁹, S(O)_nR⁹, NR¹⁰R¹

 $C(Y)OR^{11}$ and $C(Y)NR^{10}R^{11}$;

R⁸ is selected from the group consisting of hydrogen, F, Cl, Br, I, C₁-C₈ alkyl, C₁-C₈ haloalkyl, C₁-C₈ heteroalkyl, aryl, heteroaryl, OR⁹, S(O)_nR⁹, NR¹⁰R¹¹, C(Y)OR¹¹ and $C(Y)NR^{10}R^{11}$;

R⁹ is selected from the group consisting of hydrogen, C₁-C₈ alkyl, C₁-C₈

haloalkyl, C₁-C₈ heteroalkyl, aryl, heteroaryl and arylalkyl;

R¹⁰ is selected from the group consisting of hydrogen, C₁-C₈ alkyl, C₁-C₈ haloalkyl, C₁-C₈ heteroalkyl, aryl, heteroaryl, arylalkyl, CO₂R¹², C(O)R¹², SO₂R¹² and $S(O)R^{12}$;

R¹¹ and R¹² each independently is selected from the group consisting of hydrogen, C₁-C₈ alkyl, C₁-C₈ haloalkyl, C₁-C₈ heteroalkyl, aryl, heteroaryl, arylalkyl;

Page 5 of 11

PATENT NO.

: 7,214,690 B2

APPLICATION NO.: 10/080503

DATED

: May 8, 2007

INVENTOR(S)

: Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

 R^{13} is selected from the group consisting of C_1 – C_8 alkyl, C_1 – C_8 haloalkyl, C₁-C₈ heteroalkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₃-C₈ cycloalkyl, aryl, heteroaryl, arylalkyl and heteroarylalkyl;

m is selected from the group consisting of 0, 1 and 2;

n is selected from the group consisting of 0, 1 and 2;

W is selected from the group consisting of NH, $N\{R^{13}\}$, $N\{C(Y)R^{11}\}$ and $N{SO_2R^{11}};$

X is O;

Z is selected from the group consisting of NH, $N\{R^{11}\}$, $N\{C(Y)R^{11}\}$,

 $N{SO_2R^{12}}$ and $N{S(O)R^{12}}$; and

Y is O;
Column 83, Inves 53-15 and pharmaceutically acceptable salts thereof.
Claim 24 Should read and pharmaceutically acceptable salts thereof.

24. A compound according to claim 23, wherein R⁹ is selected from the group

consisting of hydrogen and optionally substituted C1-C4 alkyl.

40. A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a compound of formula:

John 85, line 20 thru column 86, line 64

should read

Z

wherein:

R¹ is selected from the group consisting of hydrogen, F, Cl, Br, I, NO₂, OR⁹, $NR^{10}R^{11}$, $S(O)_nR^9$, optionally substituted C_1 – C_8 alkyl, optionally substituted C_1 – C_8 haloalkyl, optionally substituted C_1 – C_8 heteroalkyl, optionally substituted C_3 – C_8 cycloalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted C2-C8 alkynyl and optionally substituted C_2 – C_8 alkenyl;

R² is selected from the group consisting of hydrogen, F, Cl, Br, I, CF₃, CF₂Cl, CF₂H, CFH₂, CF₂OR⁹, CH₂OR⁹, OR⁹, S(O)_nR⁹, NR¹⁰R¹¹, optionally substituted C₁–C₈ alkyl, optionally substituted C_1 – C_8 haloalkyl, optionally substituted C_1 – C_8 heteroalkyl, optionally substituted C3-C8 cycloalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted C2-C8 alkynyl and optionally substituted C2-C8 alkenyl;

PATENT NO.

: 7,214,690 B2

Page 6 of 11

APPLICATION NO.: 10/080503

DATED

: May 8, 2007

INVENTOR(S)

: Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

 R^3 and R^4 each independently is selected from the group consisting of hydrogen, $OR^9,\,S(O)_nR^9,\,NR^{10}R^{11},\,C(Y)OR^{11},\,C(Y)NR^{10}R^{11},$ optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted C3-C8 cycloalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted C₂-C₈ alkynyl and optionally substituted C₂-C₈ alkenyl;

R⁵ and R⁶ each independently are selected from the group consisting of hydrogen, CF₃, CF₂Cl, CF₂H, CFH₂, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted C₃-C₈ cycloalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted C2-C8 alkynyl and optionally substituted C2-C8 alkenyl;

R⁷ is selected from the group consisting of hydrogen, F, Cl, Br, I, optionally substituted C1-C8 alkyl, optionally substituted C1-C8 haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl, OR⁹, $S(O)_n R^9$, $NR^{10}R^{11}$, $C(Y)OR^{11}$ and $C(Y)NR^{10}R^{11}$;

R⁸ is selected from the group consisting of hydrogen, F, Cl, Br, I, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl, OR⁹, $S(O)_n R^9$, $NR^{10}R^{11}$, $C(Y)OR^{11}$ and $C(Y)NR^{10}R^{11}$

R9 is selected from the group consisting of hydrogen, optionally substituted C_1 – C_8 alkyl, optionally substituted C_1 – C_8 haloalkyl, optionally substituted C_1 – C_8 heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl and optionally substituted arylalkyl;

R¹⁰ is selected from the group consisting of hydrogen, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted arylalkyl, CO₂R¹², C(O)R¹², SO₂R¹² and S(O)R¹²;

R¹¹ and R¹² each independently is selected from the group consisting of hydrogen, optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted aryl, optionally substituted heteroaryl and optionally substituted arylalkyl;

R¹³ is selected from the group consisting of optionally substituted C₁-C₈ alkyl, optionally substituted C₁-C₈ haloalkyl, optionally substituted C₁-C₈ heteroalkyl, optionally substituted C₂-C₈ alkenyl, optionally substituted C₂-C₈ alkynyl, optionally substituted C₃-C₈ cycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted arylalkyl and optionally substituted heteroarylalkyl;

m is 1:

n is selected from the group consisting of 0, 1 and 2;

Page 7 of 11

PATENT NO.

: 7,214,690 B2

APPLICATION NO.: 10/080503

DATED

: May 8, 2007

INVENTOR(S)

: Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

W is selected from the group consisting of NH, N{R¹³}, N{C(Y)R¹¹} and $N{SO_2R^{11}};$

X is O:

Z is selected from the group consisting of NH, $N\{R^{11}\}$, $N\{C(Y)R^{11}\}$, $N{SO_2R^{12}}$ and $N{S(O)R^{12}}$; and

Y is O:

and pharmaceutically acceptable salts thereof; wherein:

the substituents of an optionally substituted group comprise one or more substituents independently selected from among alkyl, alkenyl, alkynyl, heteroalkyl, haloalkyl, haloalkenyl, haloalkynyl, cycloalkyl, aryl, heteroaryl, arylalkyl, heteroarylalkyl, alkoxy, aryloxy, haloalkoxy, amino, alkylamino, dialkylamino, alkythio, arylthio, heteroarylthio, oxo, carboxyester, carboxamido, acyloxy, hydrogen, F, Cl, Br, I, CN, NO₂, NH₂, N₃, NHCH₃, N(CH₃)₂, SH, SCH₃, OH, OCH₃, OCF₃, CH₃, CF₃, C(O)CH₃, CO₂CH₃, CO₂H, C(O)NH₂, OR⁹, SR⁹, NR¹⁰R¹¹, CF₂CF₃, CH₂CH₂F and CH₂CF₃. Column 86, line 27 thru Column 90, line 35, Claim 57 should read 57. A compound selected from the group consisting of:

(3R)-2,3,4,7-Tetrahydro-3-methyl-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f] -quinolin-8-one;

(3R)-2,3,4,7-Tetrahydro-3,4-dimethyl-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one:

(3R)-4-Ethyl-2,3,4,7-tetrahydro-3-methyl-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one;

(3R)-2,3,4,7-Tetrahydro-3-methyl-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl) -8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-2,3,4,7-Tetrahydro-3-methyl-4-propyl-10-(trifluoromethyl)-8H-[1,4] oxazino[2,3-f]-quinolin-8-one:

(3R)-4-Allyl-2,3,4,7-tetrahydro-3-methyl-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one:

(3R)-3-Ethyl-2,3,4,7-tetrahydro-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f] quinolin-8-one;

(3R)-3-Ethyl-2,3,4,7-tetrahydro-4-methyl-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one;

(3R)-3,4-Diethyl-2,3,4,7-tetrahydro-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one:

(3*R*)-3-Ethyl-2,3,4,7-tetrahydro-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl) -8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-4-(2-Chloro-2,2-difluoroethyl)-3-ethyl-2,3,4,7-tetrahydro -10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

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: 7,214,690 B2

Page 8 of 11

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It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

(3R)-4-(2,2-Difluoroethyl)-3-ethyl-2,3,4,7-tetrahydro-10-(trifluoromethyl)-8H -[1,4]oxazino[2,3-f]quinolin-8-one:

(3R)-3-Ethyl-2,3,4,7-tetrahydro-4-propyl-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one;

(3R)-4-Allyl-3-ethyl-2,3,4,7-tetrahydro-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one;

(3R)-3-Ethyl-2,3,4,7-tetraydro-4-isobutyl-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one;

(3R/S)-2,3,4,7-Tetrahydro-3-propyl-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one:

(3R/S)-2,3,4,7-Tetrahydro-4-methyl-3-propyl-10-(trifluoromethyl)-8H[1,4] oxazino-[2,3-f]quinolin-8-one;

(3R/S)-4-Ethyl-2,3,4,7-tetrahydro-3-propyl-4-(2,2,2-trifluoroethyl)-10 -(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R/S)-2,3,4,7-Tetrahydro-3-propyl-4-(2,2,2-trifluoroethyl)-10 -(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-2,3,4,7-Tetrahydro-3-isopropyl-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]-quinolin-8-one;

(3R)-2,3,4,7-Tetrahydro-3-isopropyl-4-methyl-10-(trifluoromethyl)-8H-[1,4] oxazino-[2,3-f]quinolin-8-one;

(3R)-4-Ethyl-2,3,4,7-tetrahydro-3-isopropyl-10-(trifluoromethyl)-8H-[1,4] oxazino-[2,3-f]quinolin-8-one;

(3R)-2,3,4,7-Tetrahydro-3-isopropyl-4-(2,2,2-trifluoroethyl)-10 -(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-4-(2-Chloro-2,2-difluoroethyl)-2,3,4,7-tetrahydro-3-isopropyl-10 -(trifluoromethyl)-8*H*-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-4-(2,2-Difluoroethyl)-2,3,4,7-tetrahydro-3-isopropyl-10-(trifluoromethyl) -8*H*-[1,4]oxazino[2,3-*f*]quinolin-8-one;

(3R)-4-Allyl-2,3,4,7-tetrahydro-3-isopropyl-10-(trifluoromethyl)-8H-[1,4] oxazino-[2,3-f]quinolin-8-one;

(3R)-2,3,4,7-Tetrahydro-3-phenyl-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f] quinolin-8-one:

(3*R*)-2,3,4,7-Tetrahydro-3-phenyl-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl) -8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3*R*)-4-Cyclopropylmethyl-2,3,4,7-tetrahydro-3-phenyl-10-(trifluoromethyl) -8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3*R*)-3-Benzyl-2,3,4,7-tetrahydro-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl) -8H-[1,4]oxazino[2,3-f]quinolin-8-one;

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INVENTOR(S)

: Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

2,3,4,7-Tetrahydro-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one; 2,3,4,7-tetrahydro-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl)-8H-[1,4]oxazino [2,3-f]quinolin-8-one;

Page 9 of 11

(7aR,10aS)-7,7a,8,9,10,10a-Hexahydro-1-(trifluoromethyl)

-7-(2,2,2-trifluoroethyl)-4H-cyclopenta[5,6][1,4]oxazino[2,3-f]quinolin-3-one; (7aR, 10aS)-7-Ethyl-7,7a,8,9,10,10a-hexahydro-1-(trifluoromethyl)

-4H-cyclopenta-[5,6][1,4]oxazino[2,3-f]quinolin-3-one;

(7aR, 10aS)-7,7a,8,9,10,10a-Hexahydro-3-isopropoxy-1-(trifluoromethyl)

-7-(2,2,2-trifluoroethyl)-4H-cyclopenta[5,6][1,4]oxazino[2,3-f]quinolin-3-one; (\pm) -(2S,3R)-2,3,4,7-Tetrahydro-2,3-dimethyl-4-(2,2,2-trifluoroethyl)

-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one:

(6aR)-6a,7,8,9-Tetrahydro-4-(trifluoromethyl)-1H,6H-pyrrolo[1',2':4,5][1,4] -oxazino[2,3-f]quinolin-2-one:

2,3,4,7-Tetrahydro-2,2,4-trimethyl-10-(trifluoromethyl)-8H-[1,4]oxazino

[2,3-f]-quinolin-8-one: (3*R*)-8-Chloro-3-ethyl-3,4-dihydro-8-isopropoxy-4-(2,2,2-trifluoroethyl)

-10-(trifluoromethyl)-2H-[1,4]oxazino[2,3-f]quinoline;

(3R)-3-Ethyl-3,4-dihydro-8-isopropoxy-8-methoxy-4-(2,2,2-trifluoroethyl)

-10-(trifluoromethyl)-2*H*-[1,4]oxazino[2,3-f]quinoline:

(±)-2,3,4,7-Tetrahydro-4-(2,2,2-trifluoroethyl)-3,10-bis(trifluoromethyl)

-8*H*-[1,4]oxazino[2,3-*f*]quinolin-8-one;

(-)-2,3,4,7-Tetrahydro-4-(2,2,2-trifluoroethyl)-3,10-bis(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one:

(+)-2,3,4,7-Tetrahydro-4-(2,2,2-trifluoroethyl)-3,10-bis(trifluoromethyl)

-8*H*-[1,4]oxazino[2,3-*f*]quinolin-8-one;

 (\pm) -2,3,4,7-Tetrahydro-3-(2,2,2-trifluoroethyl)-10-(trifluoromethyl)-8H-[1,4] oxazino[2,3-f]quinolin-8-one;

(±)-2,3,4,7-Tetrahydro-4-methyl-3-(2,2,2-trifluoroethyl)-10-(trifluoromethyl) -8H-[1,4]oxazino[2,3-f]quinolin-8-one:

(±)-4-Ethyl-2,3,4,7-tetrahydro-3-(2,2,2-trifluoroethyl)-10-(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(±)-2,3,4,7-Tetrahydro-3,4-bis(2,2,2-trifluoroethyl)-10-(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one:

(-)-2,3,4,7-Tetrahydro-3,4-bis(2,2,2-trifluoroethyl)-10-(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(+)-2,3,4,7-Tetrahydro-3,4-bis(2,2,2-trifluoroethyl)-10-(trifluoromethyl -8*H*-[1,4]oxazino[2,3-*f*]quinolin-8-one;

PATENT NO.

: 7.214,690 B2

APPLICATION NO.: 10/080503

DATED

: May 8, 2007

INVENTOR(S)

: Higuchi et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

(±)-4-Cyclopropylmethyl-2,3,4,7-tetrahydro-3-(2,2,2-trifluoroethyl)

-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3*R*)-4-Cyclopropylmethyl-3-ethyl-2,3,4,7-tetrahydro-10-(trifluoromethyl).

Page 10 of 11

-8H-[1,4] oxazino [2,3-f] quinolin-8-one;

(3*R*)-4-(2-Chloroethyl)-2,3,4,7-tetrahydro-3-isopropyl-10-(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(±)-2,3,4,7-Tetrahydro-2-methyl-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one:

(3R)-3-Ethyl-4-(2-hydroxy-2-methylpropyl)-2,3,4,7-tetrahydro

-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one; and

(3R)-2,3,4,7-Tetrahydro-3-isobutyl-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one; and

Column 90, line 35 pharmaceutically acceptable salts thereof.

(3R)-2,3,4,7-Tetrahydro-3-methyl-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl)

(\sim 58 -8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-3-Ethyl-2,3,4,7-tetrahydro-4-(2,2,2-trifluoroethyl)-10-(trifluoromethyl) Shovid read

-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-4-(2-Chloro-2,2-difluoroethyl)-3-ethyl-2,3,4,7-tetrahydro

-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-4-(2,2-Difluoroethyl)-3-ethyl-2,3,4,7-tetrahydro-10-(trifluoromethyl)

-8*H*-[1,4]oxazino[2,3-*f*]quinolin-8-one;

(3R)-2,3,4,7-Tetrahydro-3-isopropyl-4-(2,2,2-trifluoroethyl)

-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(3R)-4-(2-Chloro-2,2-difluoroethyl)-2,3,4,7-tetrahydro-3-isopropyl

-10-(trifluoromethyl)-8*H*-[1,4]oxazino[2,3-*f*]quinolin-8-one;

(3R)-4-(2,2-Difluoroethyl)-2,3,4,7-tetrahydro-3-isopropyl-10-(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(7aR,10aS)-7-Ethyl-7,7a,8,9,10,10a-hexahydro-1-(trifluoromethyl

-4H-cyclopenta[5,6][1,4]oxazino[2,3-f]quinolin-3-one;

(7aR,10aS)-7-7a,8,9,10,10a-Hexahydro-1-(trifluoromethyl)

-7-(2,2,2-trifluoroethyl)-4H-cyclopenta[5,6][1,4]oxazino[2,3-f]quinolin-3-one;

 (\pm) -(2S,3R)-2,3,4,7-Tetrahydro-2,3-dimethyl-4-(2,2,2-trifluoroethyl)

-10-(trifluoromethyl)-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

(±)-2,3,4,7-Tetrahydro-4-(2,2,2-trifluoroethyl)-3,10-bis(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one;

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: 7,214,690 B2

Page 11 of 11

APPLICATION NO.: 10/080503

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: May 8, 2007

INVENTOR(S)

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It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

(-)-2,3,4,7-Tetrahydro-4-(2,2,2-trifluoroethyl)-3,10-bis(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-8-one; and

(+)-2,3,4,7-Tetrahydro-4-(2,2,2-trifluoroethyl)-3,10-bis(trifluoromethyl)

-8H-[1,4]oxazino[2,3-f]quinolin-one, and

pharmaceutically acceptable salts thereof.

This certificate supersedes certificate of correction issued June 17, 2008,